In the Claims:

- 1. (currently amended) Orally An orally administrable film-shaped medicament containing at least one of the active substance deoxypeganine or/and and a deoxypeganine derivative.
- 2. (currently amended) Medicament The medicament according to claim 1, eharacterized in that it wherein said medicament contains at least one of a pharmaceutically acceptable salt of deoxypeganine or/and and a pharmaceutically acceptable salt of a derivative of deoxypeganine, with deoxypeganine hydrochloride and deoxypeganine hydrobromide being preferred as salts.
- 3. (currently amended) Medicament The medicament according to claim claims 1, wherein said medicament or 2, characterized in that it is suitable for transmucosal, especially buccal, administration of said at least one [[the]] active substance substance(s) contained therein.
- 4. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it has at least one polymer-containing layer which serves as an active substance reservoir and contains [[the]] said at least one active substance(s) substance, and wherein said with the polymer portion is present in an amount between amounting to 10 to 90%-wt, preferably 20 to 70%-wt, particularly preferably 20 to 60%-wt.
- 5. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it has a two-, three- or multilayer structure, [[with]] wherein at least one layer contains containing an active substance selected from the group comprising consisting of deoxypeganine, deoxypeganine derivatives and salts of the said substances deoxypeganine and deoxypeganine derivatives.
- 6. (currently amended) Medicament The medicament according to claim 1, wherein any one of the preceding claims, characterized in that the content of said at least one active substance content is 0.5 to 40%-wt, preferably 5 to 30% wt.
- 7. (currently amended) Medicament The medicament according to claim 1, wherein the any one of the preceding claims, characterized in that its overall thickness of said medicament is 0.05 to 3 mm, preferably 0.1 to 1 mm, especially preferably 0.1 to 0.5 mm.

- 8. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it is mucoadhesive or has at least one mucoadhesive outer surface.
- 9. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it is soluble in an aqueous media, especially in saliva, and wherein it being preferred that the dissolution takes place within between 1 second [[s up to]] and 5 minutes min, especially preferably within 3 to 30 s.
- 10. (currently amended) Medicament The medicament according to claim 1, wherein said medicament is any one of the preceding claims, characterized in that it quickly disintegrates disintegratable in an aqueous media, especially in saliva, and wherein the disintegration takes place preferably within 1 [[s]] second to 5 minutes min, especially preferably within 3 to 30 s.
- 11. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it is capable of gelatinizing gelatinizable or swelling swellable in an aqueous media, especially in saliva.
- 12. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it has a depot effect or releases [[the]] said at least one active substance substance(s) with a delay in time, preferably over a period of time of up to 8 hours h, especially up to 24 h.
- 13. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it has at least one rapidly releasing active substance-containing layer and at least one layer with retarded active substance release.
- 14. (currently amended) Medicament The medicament according to claim 1, wherein said medicament any one of the preceding claims, characterized in that it additionally contains at least one further pharmaceutically active substance which is not selected from the group including consisting of deoxypeganine, deoxypeganine derivatives and salts of deoxypeganine and deoxypeganine derivatives the said substances.
- 15. (currently amended) Film-shaped The film-shaped medicament according to claim

 1, wherein said medicament any one of the preceding claims, characterized in

that it contains at least one or more auxiliary substances substance.

- 16. (currently amended) [[Use]] A use of at least one cholinergic active substance acting on the central nervous system selected from the at least one active substance recited substances mentioned in claim claims 1 [[and 2]] for the production of producing an oral, film-shaped medicament for administering [[the]] said at least one active substance substance(s) for the treatment of treating diseases or symptoms caused by acetylcholine deficiency or where such a deficiency occurs, as well as for the treatment of treating at least one of diseases where a deficiency of endogenous amine occurs and/or and diseases which can be favourably influenced by inhibition of monoaminoxidase, said use comprising the step of introducing said medicament into the oral cavity at intervals between 1 and 6 hours and wherein the daily dose of said at least one active substance is between 50 and 750 mg.
- 17. (currently amended) [[Use]] The use according to claim 16, characterized in that the wherein said film-shaped medicament is a medicament according to one of the claims claim 1 [[to 15]].
- 18. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for the treatment of treating Alzheimer's disease or [[of]] symptoms caused by Alzheimer's disease.
- 19. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for the treatment of treating a condition selected from the group of conditions consisting of depressions, schizophrenia [[or]] and manic disorders.
- 20. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for treating chronic fatigue syndrome or disturbed sleep.
- 21. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for treating abuse of alcohol or for treating abuse of nicotine.
- 22. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for the therapy of abuse of chemical substances, especially psychotropic substances, or the dependence on such substances.
- 23. (currently amended) [[Use]] The use according to claim 16, wherein or 17,

- characterized in that the medicament is used for the prophylactic treatment of poisonings caused by organophosphorous cholinesterase inhibitors.
- 24. (currently amended) [[Use]] The use according to claim 16, wherein or 17, characterized in that the medicament is used for the treatment of treating disorders of the central nervous system, particularly impaired memory, which have been caused by the action of psychotropic substances.
- 25. (new) The medicament according to claim 2, wherein said pharmaceutically acceptable salt of a derivative of deoxypeganine is selected from the group consisting of deoxypeganine hydrochloride and deoxypeganine hydrobromide.
- 26. (new) The medicament according to claim 3, wherein said medicament is suitable for buccal, administration of said at least one active substance.
- 27. (new) The medicament according to claim 4, wherein said polymer portion is present in an amount between 20 to 70%-wt.
- 28. (new) The medicament according to claim 27, wherein said polymer portion is present in an amount between 20 to 60%-wt.
- 29. (new) The medicament according to claim 6, wherein the content of said at least one active substance is 5 to 30%-wt.
- 30. (new) The medicament according to claim 7, wherein the overall thickness of said medicament is 0.1 to 1 mm.
- 31. (new) The medicament according to claim 30, wherein the overall thickness of said medicament is 0.1 to 0.5 mm.
- 32. (new) The medicament according to claim 9, wherein said aqueous media is saliva.
- 33. (new) The medicament according to claim 9, wherein said dissolution takes place between 3 and 30 seconds.
- 34. (new) The medicament according to claim 10, wherein said aqueous media is saliva.
- 35. (new) The medicament according to claim 10, wherein said disintegration takes place within 3 to 30 seconds.
- 36. (new) The medicament according to claim 11, wherein said aqueous media is saliva.
- 37. (new) The medicament according to claim 12, wherein said delay in time is a period up to 24 hours.
- 38. (new) The use according to claim 24, wherein the medicament is used for treating disorders of impaired memory.